07-17-00 A

ATTORNEY DOCKET NO.: P- 4824.06 con
Express Mail EL084630860US

PATENT
Total Pages _____

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE
UTILITY PATENT APPLICATION TRANSMITTAL

jc682 U.S. PTO 07/14/00

FIRST NAMED INVENTOR OR APPLICATION IDENTIFIER: **DAVID D. VERNESS, GEORGE M. HUEPENBECKER, DALE A. WAHLSTROM**
MEDICAL ELECTRICAL LEAD

CERTIFICATE UNDER 37 CFR §1.10: I hereby certify that this Utility Patent Application Transmittal and the documents referred to as enclosed therein are being deposited with the United States Postal Service, in an envelope addressed to: Box Patent Application, Assistant Commissioner of Patents, Washington, D.C. 20231, "EXPRESS No. EL084630860, on this 14th day of July, 2000.

FRAYDA M. NITSCHKE
Printed Name

Frayda M. Nitschke
Signature

jc682 U.S. PTO 09/616592 07/14/00

Assistant Commissioner for Patents
BOX PATENT APPLICATION
Commissioner of Patents and Trademarks
Washington, D.C. 20231

Sir:

We are transmitting herewith the attached:

X **Patent Application Transmittal**

X **Specification:**
Total pages: 20 (including claims and abstract:Spec. 15 sheets; Claims 4 sheets; Abstract - 1

X **Drawings:**
Total sheets: 10
☐ formal X informal

X **Combined Declaration and Power of Attorney:**
☐ newly executed
X copy from prior application
☐ Deletion of Inventor(s) - Signed statement attached deleting inventor(s) named in the prior application (37 CFR 1.63(d)(2) and 1.33(b)
X Incorporation by Reference - *The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied above is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein.*

X **Accompanying application parts:**
X Notification of filing a CONTINUATION APPLICATION
X ASSOCIATE POWER OF ATTORNEY
☐ Assignment of the Invention to Medtronic, Inc.
☐ Assignment cover sheet
☐ Information Disclosure Statement
☐ PTO Form 1449
☐ Copies of IDS citations
☐ Preliminary Amendment
☐ A copy of the Petition or Conditional Petition for Extension of Time in the prior application.
X Return Postcard

IF A CONTINUING APPLICATION:

X **Continuation** ☐ **Divisional** ☐ **Continuation-in-part (CIP)**
of prior application No. 09 / 482,775 FILED JANUARY 13, 2000 WHICH IS A DIVISIONAL APPLICATION OF U.S. PATENT NO. 6,061,598 GRANTED MAY 9, 2000, WHICH IS A DIVISIONAL OF U.S. PATENT 6,018,683 GRANTED JANUARY 25, 2000, WHICH IS A DIVISIONAL APPLICATION OF APPLICATION 08/843,763 FILED APRIL 21, 1997.

X **Amend the specification by inserting before the first line the sentence: This application is a X CONTINUATION**
of prior application No. 09 / 482,775 FILED JANUARY 13, 2000 WHICH IS A DIVISIONAL APPLICATION OF U.S. PATENT NO. 6,061,598 GRANTED MAY 9, 2000, WHICH IS A DIVISIONAL OF U.S. PATENT 6,018,683 GRANTED JANUARY 25, 2000, WHICH IS A DIVISIONAL APPLICATION OF APPLICATION 08/843,763 FILED APRIL 21, 1997.

☐ **Cancel in this application original claims _____of the prior application before calculating the filing fee. (At least the original independent claim must be retained for filing purposes.)**

X **The prior application is assigned of record to Medtronic, Inc.**

X **The Power of Attorney in the prior application is to: GIRMA WOLDE-MICHAEL.**

☐ This application claims the benefit of U.S. Provisional Application(s) Serial No.(s) ______, filed _________.

X Address all future correspondence to:

GIRMA WOLDE-MICHAEL, Reg. No. 36,724
Medtronic, Inc., MS 301
7000 Central Avenue NE
Minneapolis, Minnesota 55432
phone: (763)514-6402

FEE CALCULATION	No. of Claims Filed	Claims Included in Base Fee	No. of Extra Claims	Rate	Fee
Total Claims	37	20 =	17	x 18	306
Independent Claims	1	3 =	0	x 78	0
Multiple Dependent Claims	YES			+ 260	260
Basic Filing Fee					$ 760
				TOTAL	**1,326**

X Charge Deposit Account No. 13-2546 the sum of $1,326.00 (Filing Fee) for a total of **$1,326.00.**

X The Commissioner is hereby authorized to charge any fees which may be required under 37 CFR 1.16 and 1.17, or credit any overpayment to Deposit Account No. 13-2546.. A duplicate of this transmittal is enclosed.

07/14/2000
Date

[signature]

GIRMA WOLDE-MICHAEL Reg. No.36,724
MEDTRONIC, INC.
7000 Central Avenue N.E.
Minneapolis, Minnesota 55432
Telephone: (763) 514- 6402

PATENT
ATTORNEY DOCKET: P- 4024

APPLICATION FOR UNITED STATES LETTERS PATENT
for

MEDICAL ELECTRICAL LEAD

by

David D. Verness
George M. Huepenbecker
Dale A. Wahlstrom

ATTORNEY OF RECORD:

Reed A. Duthler, Reg. No. 30,626
MEDTRONIC, INC.
7000 Central Avenue N.E
Minneapolis, Minnesota 55432
Telephone: (612) 514- 3351
Facsimile: (612) 514-3233

CERTIFICATE OF "EXPRESS MAIL"

Mailing Label No. EM 575726767US

Date of Deposit: April 21, 1997

I hereby certify that this paper or fee is being deposited with the United States Postal Service as "EXPRESS MAIL" POST OFFICE TO ADDRESSEE" service under 37 CFR 1.10 on the date indicated above and is addressed to the Commissioner of Patents and Trademarks, Washington, D.C. 20231.

Printed Name Frayda M. Nitschke

Signature Frayda M. Nitschke

Docket: P-4024

MEDICAL ELECTRICAL LEAD

Background of the Invention

The present invention relates to implantable electrical leads generally, and more specifically to cardiac pacing leads.

The conductors in cardiac pacing leads occasionally have a tendency to fracture due to repetitive application of stress to the conductor. One way in which this problem has previously been addressed is by reinforcing the lead body in the area in which stress is to be expected, as in U.S. Patent No. 5,545,203, issued to Doan et al. This patent is directed primarily toward reinforcing the lead against fracture due to application of compressive forces. Reinforcement of the lead body is also disclosed in U.S. Patent No. 5,591,142, issued to Van Erp et al.. It has also been proposed to reinforce the lead body by means of adding a tensile reinforcement as in U.S. Patent No. 5,231,996 issued to Bardy et al. In this patent, the lead is provided with a non-conductive tensile member such as a polyester cord, which runs the length of the lead body. Other leads having cords or reinforcements running throughout their length are disclosed in U.S. Patent No. 3,844,292 and U.S. Patent No. 3,572,344 issued to Bolduc. A third proposal for dealing with the possibility of conductor fracture is to render the portion of the lead body in direct contact with the conductor conductive by addition of carbon or other conductive material, as disclosed in U.S. Patent No. 4,033,355, issued to Ammundson.

Summary of the Invention

The present invention is directed toward providing a lead which has an increased resistance to fracture and has the capability of continued function after fracture of a conductor. The lead is provided with a coiled conductor which may be monofilar or multifilar and which extends along the length of the lead, running from an electrical connector at the proximal end of the lead to an electrode at or near the distal end of the lead. In addition, the lead is provided with a stranded conductor

which extends along the coiled conductor from a point along the lead body located proximal to the point of expected breakage of the coiled conductor to a point along the lead body located distal to the point of expected breakage. The proximal and distal ends of the stranded conductor in some embodiments are electrically and mechanically coupled to the coiled conductor, limiting the extensibility of the coiled conductor, rendering the coiled conductor less susceptible to axially applied tensile forces and also providing for continued electrical connection to the electrode, in the event that the coiled conductor fractures intermediate the proximal and distal ends of the stranded conductor. In alternative embodiments, the stranded conductor may be coupled only at its proximal or distal end to the coiled conductor or may simply be located in the same lumen of the lead as the coiled conductor, without mechanical connection to the coiled conductor.

Brief Description of the Drawings

Figure 1 is a plan view of an implantable lead in which the present invention is practiced.

Figure 2 is cross-sectional view through the lead of Figure 1, illustrating a first embodiment of the invention.

Figure 3 is cross-sectional view through the lead of Figure 1, illustrating a second embodiment of the invention.

Figure 4 is a side, cut-away view through the lead of Figure 1, illustrating the first embodiment of the invention..

Figure 5 is a side, cut-away view through the lead of Figure 1, illustrating the second embodiment of the invention.

Figure 6 is a side, cut-away view through the distal portion of the lead of Figure 1, illustrating the first embodiment of the invention..

Figure 7 is a side, cut-away view through the distal portion of the lead of Figure 1, illustrating the second embodiment of the invention.

Figure 8 is a side, cut-away view through the connector assembly of the lead of Figure 1, illustrating the first embodiment of the invention..

Figure 9 is a side, cut-away view through the connector assembly of the lead of Figure 1, illustrating the second embodiment of the invention.

Figure 10 is a side, cut-away view through the lead of Figure 1, illustrating a third embodiment of the invention.

Figure 11 is cross-sectional view through the lead of Figure 1, illustrating a fourth embodiment of the invention.

Figure 12 is a side, cut-away view through the lead of Figure 1, illustrating the fourth embodiment of the invention..

Figure 13 is cross-sectional view through the lead of Figure 1, illustrating a fifth embodiment of the invention.

Figure 14 is a side, cut-away view through the lead of Figure 1, illustrating the fifth embodiment of the invention.

Figure 15 is a side, cut-away view through the a lead according to the present invention, illustrating an alternative mechanism for interconnecting a coiled conductor with a stranded conductor.

Figure 16 is a plan view of a lead having a rotatable fixation helix, embodying the present invention.

Figure 17 is a cross-sectional view through the lead of Figure 16.

Figure 18 is a side, cut-away view through the distal portion of the lead of Figure 16.

Figure 19 is a side, cut-away view through the proximal portion of the lead of Figure 16.

Detailed Description of the Preferred Embodiments

Figure 1 is a plan view of a defibrillation lead of a lead in which the present invention is practiced. The present invention, of course, may also be usefully practiced in the context of other types of medical electrical leads, such as cardiac pacing leads, nerve and muscle stimulation leads, and so forth.

The lead of Figure 1 is provided with an elongated insulative lead body 10, preferably fabricated of silicone rubber, polyurethane or other biocompatible elastomer. At the proximal end of the lead, it carries an elongated defibrillation electrode 12, a ring

electrode 14 and a tip electrode 16, each coupled to a conductor located within the lead body 10. Tines 18 are provided in maintaining electrode 16 in contact with the tissue of the right ventricle. Electrodes 16, 14 and 12 may correspond generally to conventionally available pacing and defibrillation electrodes.

The proximal end of the lead carries a connector assembly, beginning with a molded lead bifurcation 20, which splits off two of the conductors within lead body 10 to a bipolar, in-line connector assembly 24, generally corresponding to the IS-1 connector standard for pacing leads. However, other types of connector assemblies may also be adapted to practice the present invention. Connector assembly 24 is provided with a first set of sealing rings 28, a connector ring 32, a second sealing rings 34 and connector pin 36. Connector pin 36 is coupled to the conductor which extends through the lead body 10 to tip electrode 16. Connector ring is coupled to the conductor which extends through the lead body 10 to ring electrode 14. The conductor coupled to defibrillation electrode 12 extends into connector assembly 22, which carries a set of sealing rings 26 and a connector pin 36, coupled to the conductor extending through lead body 10 to defibrillation electrode 12.

In the specific context of the lead illustrated in Figure 1, the conductor coupling connector pin 36 to electrode 16 takes the form of a monofilar or multifilar coiled conductor to allow passage of a stylet therethrough, while the conductors coupling ring electrode 14 to connector ring 32 and coupling defibrillation electrode 12 to connector pin 30 take the form of bundled, stranded wires, provided with a coating of PTFE. However, the conductors coupling ring electrode 14 and defibrillation electrode 12 may take the form of any of the various conductor types known for use in conjunction with implantable electrical leads. If fewer electrodes are provided on the lead, correspondingly fewer conductors will be included. One or more physiologic sensors may be added to the lead or substituted for one or more of the illustrated electrodes. Also located within lead body 10 is a stranded wire conductor which extends along a length of the coiled conductor and which serves a mechanism for bridging a fracture of the coiled conductor which occurs between the ends of the stranded conductor. In some embodiments, the stranded conductor also couples electrode 16 to connector pin 36, providing both an axial reinforcement and a redundant electrical connection, as

described in more detail below. In other embodiments, the electrical interconnection between the coiled and stranded conductors may simply be the contact between the two conductors which occurs as a result of both conductors being located in the same lumen of the lead.

Figure 2 illustrates a cross-section through lead body 10, illustrating the inter-relation of the conductor lumens 100, 102 and 104 with compression lumens 106, 108 and 110, which are described in more detail in U.S. Patent No. 5,584,873, issued to Shoberg et al. and incorporated herein by reference in its entirety. In this view it can be seen that lumens 100 and 102 contain conductors 112 and 114 which in the illustrated embodiment may take the form of PTFE coated bundled stranded wires having a generally straight configuration. In particular, conductors 112 and 114 may take the form of a PTFE coated, bundled, stranded 49 filar cable formed of seven strands, each strand formed of seven filars, as described in more detail in U.S. Patent No. 5,584,873 by Shoberg et al. incorporated herein by reference in its entirety. Lumen 104 contains a conventional multifilar coiled conductor 116 and a small diameter bundled stranded wire conductor 118. Conductor 118 may take the form of a seven filar bundle or cable of MP35N or silver cored MP35N wire, as described in U.S. Patent No. 5,246,014, issued to Williams et al and also incorporated herein by reference in its entirety, such that conductor 118 corresponds generally to one of the seven strands that make up conductors 112 and 114. In preferred embodiments, conductor 118 may have an outer diameter of about .003 inches.

In spite of its small diameter and generally straight configuration, stranded conductor 118 is extremely resistant to fracturing in response to repeated flexure of the lead body and displays a high tensile strength. Thus, should coil conductor 116 fracture, redundant, stranded conductor 118 will remain to provide for connection to the electrode to which coiled conductor 116 is coupled. If the stranded and coiled conductors are uninsulated from one another, they make contact with one another at multiple points along the lead body, so that a break of the coiled conductor occurring between the ends occurring between the ends of the stranded conductor will be bridged. The ends of conductor 118 may also be mechanically coupled to the coiled conductor 116 and thereby serve to maintain the structural integrity of the lead, preventing partial

disassembly due to applied tensile forces. If the lead is removed, conductor 118 may thus also serve as a reinforcement, allowing traction force to be applied to the distal end of the lead during extraction. In either case, conductor 118 allows for continued functioning of the lead after fracture of the coiled conductor 116, allowing for replacement of the lead, when convenient, without interruption of the therapeutic function of the pacemaker or stimulator to which the lead is coupled.

In some embodiments of the invention, conductor 118 is uninsulated along its length and thus makes contact with conductor 116 at various points along the length of the lead. In such embodiments, it is to be expected that the conductor 118 will serve as both a redundant conductor, coupling the connector pin 36 to the electrode 16, and as a conductive bridge between the broken ends of the conductor 116, as it will be in contact with the conductor 116 on either side of the break. With this structure, changes in overall impedance between the connector pin and electrode are expected to be relatively small, allowing for essentially undiminished performance of the lead. Alternatively, conductor 118 may be provided with an insulative coating of PTFE or other insulative material. In such embodiments, conductor 118 will serve as a redundant connector, connecting connector pin 32 to electrode 16, and upon fracture of conductor 116, a substantial change in connector pin to electrode impedance will be manifested. In the context of implantable stimulators capable of monitoring changes in lead impedance, this provides the physician and/or the device itself with a mechanism for detecting the fracture in 116. However, within the context of the present invention, the fracture can be detected without the serious consequences which would otherwise be associated with disconnection of the electrode 16 from the connector pin 36. In the context of implantable stimulators having the ability to automatically adjust stimulus pulse amplitude and input amplifier sensitivity, the device may respond to the change in lead impedance by noting the occurrence of a fracture in conductor 116 and may correspondingly alter its programmed parameters in order to restore performance essentially to that preceding the fracture of conductor 116.

FIG. 3 is a cross-sectional view through an alternative embodiment of the lead illustrated in FIG. 1, in which all labeled elements correspond to identically labeled elements in FIG. 2. The embodiment illustrated in FIG. 3 differs from that illustrated in

FIG. 2 only in that stranded conductor 118 is located within the lumen of conductor 116, rather than external to conductor 116. This embodiment may be particularly advantageous in the context of leads, such as epicardial electrode leads or some nerve and muscle stimulation leads which do not require passage of a stylet through the lumen of coil conductor 116.

FIG. 4 is a side, cutaway view through the lead of FIG. 1, illustrating the first embodiment of the present invention, also illustrated in FIG. 2. In this view, it can be seen that stranded conductor 118 is loosely spiraled around coiled conductor 116 along the length of the lead, facilitating flexure of the lead body and the conductors located therein. If the ends of conductor 118 are mechanically coupled to conductor 116, this structure also allows for a limited amount of axial elongation of the lead body and conductor 116 along the length of conductor 118. All other labeled elements correspond to those illustrated in FIG. 2.

FIG. 5 shows a side cutaway view through the second embodiment of the lead of FIG. 1, also illustrated in FIG. 3. In this view, the stranded conductor is shown arranged loosely within the lumen of coiled conductor 116. All other labeled elements correspond to those illustrated in FIG. 2.

In the embodiments illustrated in Figures 2, 3, 4 and 5, conductor 118 may be insulated or uninsulated, as discussed above, depending on whether contact between the two conductors along their length is desired. An alternative embodiment in which the stranded conductor is desired to be insulated from the coiled conductor along some portion of its length may employ a separate lumen in the lead body for the stranded conductor, intermediate its points of connection to the coiled conductor. An additional alternative as discussed below may employ a tubular, insulative sheath within or around coiled conductor 116 to insulate it from conductor 118.

FIGs. 6 et seq. show basic mechanisms which may optionally be employed to mechanically interconnect the stranded conductor 118, the coiled conductor 116, electrode 16 and connector pin 36. These illustrated interconnection mechanisms are intended to be exemplary, and may of course, be employed in conjunction with other components of implantable leads, including other types of electrical connectors such as connector rings, corresponding to connector ring 32 and to interconnect these

conductors with other types of electrodes and to interconnect these components with other lead components such as physiologic sensors such as pressure sensors, oxygen sensors, temperature sensors and the like.

Figure 6 is a sectional view through the distal portion of the lead illustrated in Figure 1. In this view, the interconnection of conductor 116, conductor 118 and electrode 16 is visible. Extending distally from the defibrillation electrode 12, the lead takes the form of a molded piece part 228, which carries ring electrode 14, which is in turn coupled to stranded conductor 112 (not visible in this view). Electrode 16 as illustrated is a steroid-eluting electrode, provided with a monolithic controlled release device 222 located within a chamber within the electrode. Electrode 16 is coupled to a coiled conductor 116 and 118 by means of an external crimping sleeve 224, which compresses conductor 118 against conductor 116 and compresses conductor 116 against the proximal portion 220 of electrode 16. Other types of tip electrodes, including screw-in electrodes may of course be substituted for electrode 16. Similarly, other mechanisms may be employed to interconnect conductors 118 and 116 and electrode 16, including welding, swaging, crimping and combinations thereof, including mechanisms as disclosed in commonly assigned, copending U.S. Patent Application No. 08/657,577 by Boser et al. filed June 7, 1996 and U.S. Patent Application No. 08/439,332 by Swoyer et al., filed May 11, 1995, both incorporated herein by reference in their entireties.

Conductor 114 passes through an internal lumen 100 within lead body 10, and has its insulation removed in areas in which it passes through the cross-bore crimp sleeve 212. The distal turn of electrode coil 12 can be seen at 12A as it passes through the perpendicular cross-bore through sleeve 212. The sleeve 212 is crimped to the conductor 114 and a portion of the distal turn of electrode coil 12 is inserted through the cross bore and the entry and exit points of the coil are laser welded to the sleeve. External polymeric sleeve 230 is slid over the distal ends of conductor coil 12, and the areas between the sleeve 230 lead body 10 is backfilled by means of medical adhesive or other polymeric material.. The electrode coil 12 may be secured to the outer circumference of the lead body 10 by means of a backfilling process as described in U.S. Patent No. 4,934,049, incorporated herein by reference in its entirety.

Figure 7 illustrates the distal portion of the lead in the second embodiment of the invention in which the stranded conductor 118 is located internal to coil conductor 116. All illustrated elements correspond to identically numbered elements in Figure 6, with the exception that a bore is provided in the proximal section 220A of electrode 16, and stranded conductor 118 is crimped therein.

While Figures 6 and 7 show the inter-connection of the stranded and coiled conductors at the tip electrode 16, these conductors may instead be connected at a point proximal to the tip electrode, for example by use of a cross-bore crimp sleeve similar to sleeve 212, or by means of other types of welded, swaged or crimped connections as discussed above.

Figure 8 is sectional view through the bipolar connector assembly 24 of the lead illustrated in Figure 1, illustrating the first embodiment of the invention. In this view, the proximal end of connector pin 36 is visible in cross-section, and connector ring 32 is visible in cross-section. Connector pin 36 is coupled to coiled conductor 116 by means of a swaging core 200, which compresses conductor coil 116 and stranded conductor 118 between the interior lumen of connector pin 36 and the outer surface of swaging core 200, in a conventional fashion. An insulative sleeve 206 surrounds conductors 116 and 118, and extends distally, back through the connector assembly into molded sealing ring sleeve 28 (Fig. 1).

Surrounding connector pin 36 is a molded sealing ring sleeve 34, which may be fabricated of silicone rubber, which in turn is mounted to a spacer 204 which is typically fabricated of a harder plastic, such as polyurethane. Spacer 204 is molded in situ between connector pin 36 and ring electrode 32, and is maintained in mechanical interconnection with electrode 32 by means of internal threading 208, as described in U.S. Patent No. 4,572,605, issued to Hess, et al., incorporated herein by reference in its entirety.

Figure 9 is a sectional view through the bipolar connector assembly 24 of the lead illustrated in Figure 1, illustrating the second embodiment of the invention. All illustrated elements correspond to identically numbered elements in Figure 8, with the exception that the stranded conductor 118 is located internal to coil conductor 116.

As in the case of figures 6 and 7 above, other mechanisms may be employed to interconnect conductors 118 and 116 and connector pin 36, including welding, swaging, crimping and combinations thereof, as described above. Additionally, these conductors may instead be connected at a point distal to the connector pin, for example by use of a cross-bore crimp sleeve similar to sleeve 212, or by means of other types of welded, swaged or crimped connections as discussed above.

If it is not desired to mechanically interconnect on or both ends of the stranded conductor 118 to the coiled conductor 116, the internal structure of the leads may correspond to those illustrated in Figures 6, 7, 8 or 9 above, with the exception the stranded conductor 118 is simply not crimped, swaged or otherwise coupled to the connector pin, electrode or coiled conductor 118. In such embodiments, the stranded conductor may extend the entire length of the coiled conductor or may extend over only a portion of the length of the coiled conductor. While the Figures 6, 7, 8.and 9 illustrate the coil and stranded conductor pair coupled to the connector pin and tip electrode, it should also be understood that the invention may also be usefully practiced in leads in which these conductors are coupled to other connector elements, other electrodes and/or physiologic sensors located on the lead body. The interconnection methods of Figures 6, 7, 8 and 9 may also be used to connect the stranded conductor 118 to the coiled conductor 116 and to such other lead components.

Figure 10 illustrates a third embodiment of the invention All numbered components correspond to identically numbered components in the Figures above. In this embodiment, an uninsulated stranded conductor 118 repeatedly enters and exits the internal lumen of the coiled conductor 116, by passing between the coils. This embodiment, while more difficult to assemble, provides for in increase in the number of contact points between the stranded and coiled conductors, which may be beneficial in the case of coil fractures as it will in many case shorten the distance which the stranded conductor must bridge as compared to the first and second embodiments and may provide for more consistent contacts between the stranded and coiled conductors.

Figure 11 illustrates a cross section through a fourth embodiment of the invention All numbered components correspond to identically numbered components in the Figures above. In this embodiment the stranded conductor 118 is located outside of

coiled conductor 116 and is insulated from conductor 116 over at least a portion of its length by means of an insulative tube 300, located exterior to conductor 116. Tube 300 may be formed of PTFE or other insulative biocompatible plastic, and may extend over all or some of the length of coiled conductor 116. In this embodiment, it is desirable that the ends of stranded conductor 118 are mechanically coupled to the coiled conductor 116 on either side of the tube 300.

Figure 12 illustrates a side, cut-away view through the fourth embodiment of the invention as illustrated in Figure 11. All numbered components correspond to identically numbered components in the Figures above

Figure 13 illustrates a cross section through a fifth embodiment of the invention All numbered components correspond to identically numbered components in the Figures above. In this embodiment the stranded conductor 118 is located inside of coiled conductor 116 and is insulated from conductor 116 over at least a portion of its length by means of an insulative tube 302, located interior to conductor 116. Tube 302 may be formed of PTFE or other insulative biocompatible plastic, and may extend along all or some of the length of coiled conductor 116. In this embodiment, it is desirable that the ends of stranded conductor 118 are mechanically coupled to the coiled conductor 116 on either side of the tube 302.

Figure 14 illustrates a side, cut-away view through the fifth embodiment of the invention as illustrated in Figure 13. All numbered components correspond to identically numbered components in the Figures above.

Figure 15 illustrates an alternative mechanism for interconnecting a stranded conductor 412 with a coiled conductor 416, both located within an internal lumen of lead body 410. Conductive crimp sleeve 418 is crimped to coiled conductor 416 by crimps 420. Optionally, a cylindrical crimping core (not illustrated) may be inserted into the lumen of coiled conductor 416, prior to crimping. Stranded conductor 412 is coupled to the crimp sleeve 418 by means of conductive sleeve 422, by the following methods. Stranded conductor 412 may be threaded through sleeve 422, which is then pushed onto crimping core 418, pulling stranded conductor 412 along and compressing it between crimp sleeve 418 and sleeve 422. In conjunction with this method, the interior of sleeve 422 may be provided with threads or other internal texturing to

frictionally engage stranded conductor 412. Alternatively, stranded conductor 412 may arranged alongside crimp core 418 and sleeve 422, may then be pushed onto crimp core 418, compressing conductor 412 between crimp sleeve 418 and sleeve 422. In conjunction with this method, the exterior of crimp of sleeve 418 may be provided with threads or other external texturing to frictionally engage stranded conductor 412. As yet another alternative, sleeve 422 may simply be crimped around stranded conductor 412 and crimping sleeve 418. Crimp sleeve 418 may take the form of a portion of a connector pin or ring on the proximal end of the lead body or a portion of an electrode or other sensor on the distal portion of the lead body, or may simply be a cylindrical sleeve, employed to couple the stranded and coiled conductors at some point along the lead body. Plastic Sleeve 414 insulated stranded conductor 412 from coiled conductor 416 over a portion of their lengths.

Figure 16 is a plan view of a defibrillation lead in which the present invention is practiced, employing a tip electrode taking the form of a rotatable fixation helix 316. The lead of Figure 16 is provided with an elongated insulative lead body 310, preferably fabricated of silicone rubber, polyurethane or other biocompatible elastomer. At the distal end of the lead, it carries an elongated defibrillation electrode 312, a ring electrode 314 and a rotatable helical tip electrode 316, rotatably and advancably mounted in insulative electrode head 318. Each electrode is coupled to a conductor located within the lead body 310. Electrodes 314 and 312 may correspond generally to conventionally available pacing and defibrillation electrodes. A cap member 319 is located at the distal end of electrode head 318 and serves to retain a monolithic controlled release device as discussed below.

The proximal end of the lead carries a connector assembly, beginning with a molded lead bifurcation 320, which splits off two of the conductors within lead body 310 to a bipolar, in-line connector assembly 324, generally corresponding to the IS-1 connector standard for pacing leads. However, other types of connector assemblies may also be adapted to practice the present invention. Connector assembly 324 is provided with a first set of sealing rings 328, a connector ring 332, a second sealing rings 334 and connector pin 336. Connector pin 336 is rotatably mounted and is coupled to a rotatably mounted conductor which extends through the lead body 310 to helical electrode 316.

Connector ring 332 is coupled to a conductor which extends through the lead body 310 to ring electrode 314. A conductor coupled to defibrillation electrode 312 extends into connector assembly 322, which carries a set of sealing rings 326 and is coupled to connector pin 336.

In the specific context of the lead illustrated in Figure 16, the conductor coupling connector pin 336 to electrode 316 takes the form of a monofilar or multifilar coiled conductor to allow passage of a stylet therethrough, while the conductors coupling ring electrode 314 to connector ring 332 and coupling defibrillation electrode 312 to connector pin 330 take the form of bundled, stranded wires, provided with a coating of PTFE. However, the conductors coupling ring electrode 314 and defibrillation electrode 312 may take the form of any of the various conductor types known for use in conjunction with implantable electrical leads. If fewer electrodes are provided on the lead, correspondingly fewer conductors will be included. One or more physiologic sensors may be added to the lead or substituted for one or more of the illustrated electrodes. Also located within lead body 310 is a stranded wire conductor which extends along a length of the coiled conductor and which serves a mechanism for bridging a fracture of the coiled conductor which occurs between the ends of the stranded conductor, as discussed above.

Figure 17 illustrates a cross section through the lead illustrated in Figure 16. The lead body is provided with five lumens, including three circular lumens 350, 354 and 356 and two teardrop-shaped compression lumens 352 and 358. Coiled conductor 360 is coupled to helical electrode 316 (Fig. 16) and connector pin 336 (Fig. 16). On rotation of connector pin 336, conductor 360 transmits torque to rotate electrode 316, advancing it out the distal end of electrode head 318 (Fig. 16) and screwing it into heart tissue. Conductors 364 and 368 as illustrated are stranded or cabled conductors corresponding to conductors 112 and 114 (Fig. 2) and couple connector pin 330 to defibrillation electrode 312 and connector ring 332 to electrode 314, respectively. Stranded conductor 362 is coupled to coiled conductor 360 adjacent the proximal and distal ends of the lead, providing a redundant connector and tensile reinforcement in the same fashion as conductor 118 (Fig. 2) discussed above. The wall of lead body 310 separating lumens 350 and 352 insulates conductor 362 from conductor 360 between the

points at which they are electrically coupled. Electrical interconnection of conductors 360 and 362 is by means of rotating electrical couplings as described in conjunction with Figs. 18 and 19 below, which allow rotation of coil conductor 360 relative to stranded conductor 362.

Figure 18 is a side cut-away view through the distal portion of electrode head 318 of the lead of Fig. 16. Electrode head 318 is fabricated of a rigid, biocompatible plastic such as a polyurethane, and is provided with an internal longitudinal lumen 321. Cap 319 retains a toroidal monolithic controlled release device 374, which serves to elute an anti-inflammatory steroid such as sodium dexamethasone phosphate, as described in U.S. Patent No. 4,972,848, issued to DiDomnico and incorporated herein by reference in its entirety. Guide 363 engages helical electrode 316 such that rotation of the electrode serves to advance it out the distal end of electrode head 318 or withdraw it into lumen 321. Coiled conductor 360 is mechanically and electrically coupled to the proximal end of electrode 316 by conductive crimp sleeve 368, compressed by crimps 376. Crimp sleeve 368 is provided with a circumferential shoulder 378 which serves to limit distal movement of helix 316 by contact with radio-opaque marker ring 364 and which serves to limit proximal movement of helix 316 by contact with conductive ferrule 369.

Electrical interconnection of stranded conductor 362 and coiled conductor 360 is accomplished by ferrule 369 which is crimped to stranded conductor 362 by crimp 370 and is provided with contact means 372 for coupling to conductive crimp sleeve 368. As illustrated the contact means 372 is a conductive spring with individual turns offset from one another to springingly contact both ferrule 369 and crimp sleeve 368 while allowing rotation and longitudinal movement of crimp sleeve 368, in a manner analogous to that illustrated in U.S. patent No. 4,557,643, incorporated herein by reference in its entirety. Alternatively, coupling means in the form of other types of spring contacts, fine wire brushes or other known mechanisms for rotatable electrical couplings may be substituted.

Figure 19 shows a side, cut-away view through the lead of Figure 16 in the vicinity of bifurcation 320. In this view, coiled conductor 360 and stranded conductors 362 and 364 are visible, exiting from lead body 310 and entering into molded

bifurcation 320. Interconnection of stranded conductor 362 and coiled conductor 360 is accomplished by ferrule 380 coupled to conductor 362 by crimp 382, crimp sleeve 386 coupled to coiled conductor 360 by crimps 388 and conductive spring 384. These components function in the same way as their counterparts illustrated in Figure 18 to couple the conductors while allowing rotational movement of coiled conductor 360. As in the case of Figure 18, the known mechanisms for making a rotating electrical connection may be substituted. While the rotatable coiled conductor in this embodiment is coupled to a helical electrode, it may alternatively be coupled to any other electrode which is deployed or manipulated by applied torque and may also be employed with any other mechanism requiring both applied torque and an electrical connection.

While the embodiments described above are shown as alternatives it should be understood that they also may be combined, with the location and insulation of the stranded conductor relative to the coiled conductor varying along the length of the lead body. In addition, while the embodiments above all take the form of multiple lumen leads fabricated using multi-lumen tubing, it should be understood that the invention may also usefully be practiced in embodiments having only a single lumen and in embodiments in which multiple lumens are provided by means of single lumen tubes located within larger diameter single lumen tubes. As such, the disclosure above should be taken as exemplary, rather than limiting with regard to the scope of the claims which follow. In conjunction with the above disclosure, we claim:

Claims:

1. An implantable medical lead comprising an elongated lead body having an elongated lumen therein:

an elongated coiled conductor mounted within the lumen of the lead body;

a stranded conductor, extending along the length of the lead body and coupled electrically to the coiled conductor at a first location and a second location distal to and spaced from the first location.

2. A lead according to claim 1 wherein said lead is provided with an electrical connector located at a proximal end of the lead body and in which the coiled conductor is connected to the connector.

3. A lead according to claim 2 in which the stranded conductor and the coiled conductor are electrically and mechanically coupled to the connector.

4. A lead according to claim 1 provided with an electrode located on said lead body and in which the coiled conductor is coupled to the electrode.

5. A lead according to claim 4 wherein the coiled conductor and the stranded conductor are both electrically and mechanically coupled to the electrode.

6. A lead according to claim 1 provided with an electrical component located on said lead body, wherein said coiled conductor is coupled to said electrical component.

7. A lead according to claim 6 wherein said coiled conductor and said stranded conductor are electrically and mechanically coupled to said electrical component.

8. A lead according to claim 6 wherein said electrical component is an electrode.

9. A lead according to claim 1 wherein said stranded conductor is insulated from said coiled conductor between said first and second locations.

10. A lead according to claim 1 or claim 2 wherein said stranded conductor and said coiled conductor are both located within said lumen of said lead body.

11. A lead according to claim 10 wherein said coiled conductor defines an internal lumen and wherein said stranded conductor is located within said lumen of said coiled conductor for a distance, along the length of the lead body.

12. A lead according to claim 10 wherein said coiled conductor defines an internal lumen and said stranded conductor is located outside of said internal lumen, for distance along said lead body.

13. A lead according to claim 1 wherein said coiled conductor defines an internal lumen and wherein said stranded conductor passes between coils of said coiled conductor, to pass from within the internal lumen of the coiled conductor, to pass from within the internal lumen of the coiled conductor to outside of the coiled conductor.

14. A lead according to claim 1 wherein said stranded conductor comprises a seven stranded conductor.

15. A lead according to claim 14 wherein said stranded conductor is made of MP35N alloy.

16. A claim according to either claim 14 or 15 wherein said stranded conductor has an outer diameter of about .003 inches.

17. A lead according to claim 1 in which the stranded conductor and the coiled conductor are electrically and mechanically coupled to one another.

18. A lead according to claim 17 in which the stranded conductor and the coiled conductor are electrically and mechanically coupled to one another at said first and second locations.

19. A lead according to claim 18 in which the stranded conductor and the coiled conductor are electrically insulated from one another along a portion of the coiled conductor intermediate said first and second locations.

20. A lead according to claim 1 in which the stranded conductor and the coiled conductor are electrically insulated from one another along a portion of the coiled conductor intermediate said first and second locations.

21. A lead according to claim 20 wherein the stranded conductor and the coiled conductor are electrically insulated from one another along a portion of the coiled conductor intermediate said first and second locations by means of an insulative tubular sheath.

22 A lead according to claim 21 wherein the coiled-conductor defines an internal lumen and the tubular sheath is located within the internal lumen of the coiled conductor and the stranded conductor is located within the tubular sheath.

23 A lead according to claim 21 wherein the tubular sheath is located exterior to the coiled conductor and the stranded conductor is located exterior to the tubular sheath.

24. A lead according to claim 1 or claim 2 wherein said stranded conductor is located outside said lumen of said lead body.

25. A lead according to claim 24 wherein said lead body is provided with an additional longitudinal lumen and wherein said stranded conductor is located within said additional lumen.

26. A lead according to claim 1 wherein said coiled conductor is rotatably mounted in said lumen.

27. A lead according to claim 26 wherein said lead comprises means for coupling said stranded conductor to said coiled conductor while said coiled conductor rotates.

28. A lead according to claim 26 or claim 27 wherein said lead comprises a rotatably mounted electrical connector located on a proximal portion of said lead body and coupled to said coiled conductor.

29. A lead according to claim 28 wherein said lead comprises a rotatably mounted electrode means located on a distal portion of said electrode body and coupled to said coiled conductor.

09515592.071400

ABSTRACT OF THE DISCLOSURE

An implantable lead which has an increased resistance to fracture and has the capability of continued function after fracture of a conductor. The lead is provided with a coiled conductor which may be monofilar or multifilar and which extends along the length of the lead, running from an electrical connector at the proximal end of the lead to an electrode at or near the distal end of the lead. In addition, the lead is provided with a stranded conductor which is electrically coupled to the coiled conductor at point along the lead body located proximal to the point of expected breakage of the coiled conductor and at a point along the lead body located distal to the point of expected breakage. The proximal and distal ends of the stranded conductor in some embodiments are also mechanically coupled to the coiled conductor.

Docket No. P-4824

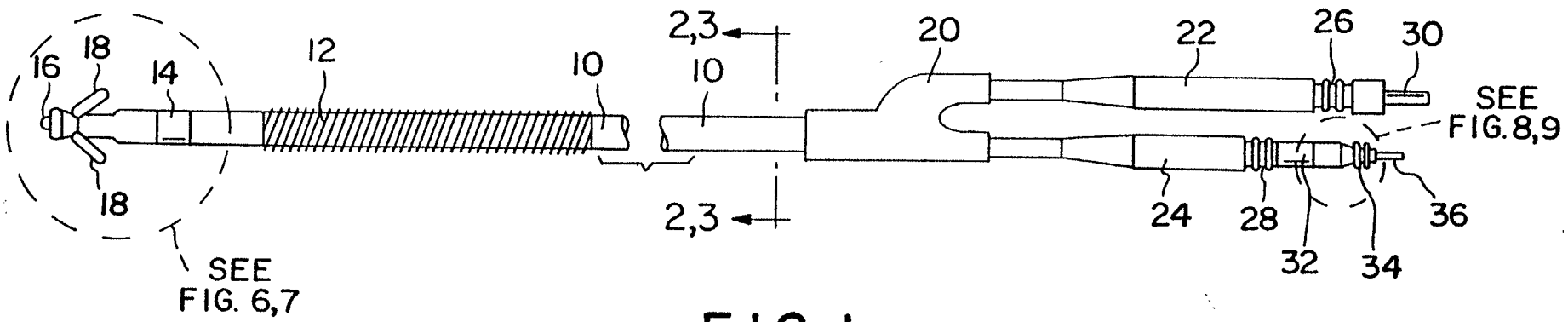
16
18
14
12
10
2,3
10
20
22
26
30
SEE
FIG. 8,9
18
2,3
24
28
32
34
36
SEE
FIG. 6,7

FIG. 1

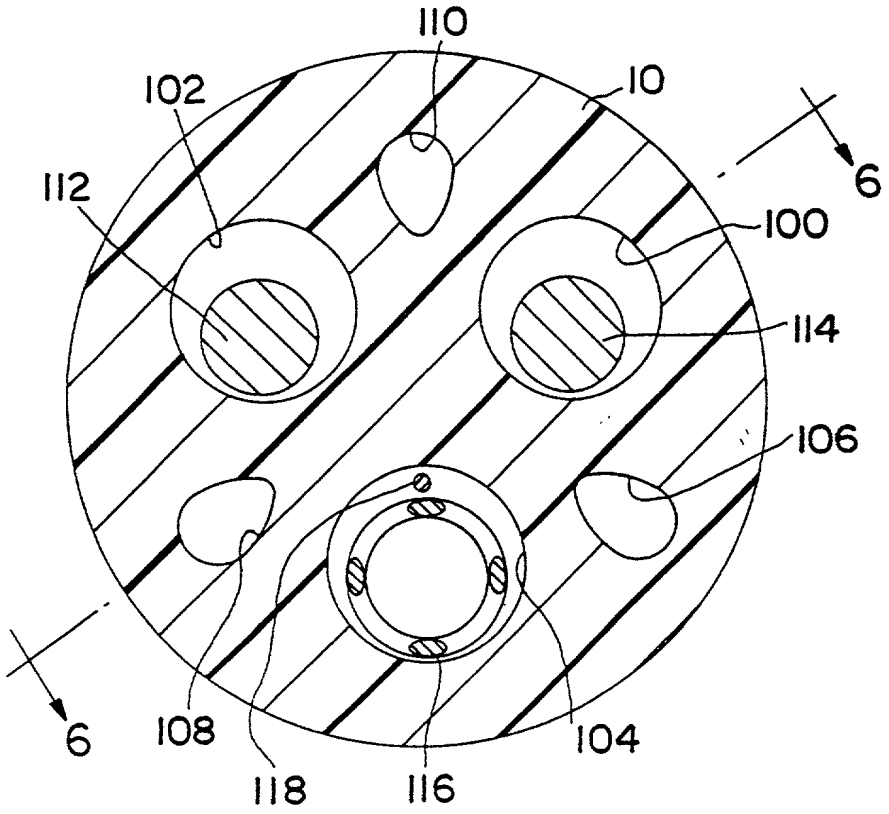
110
102
10
112
6
100
114
106
6
108
104
118
116

FIG. 2

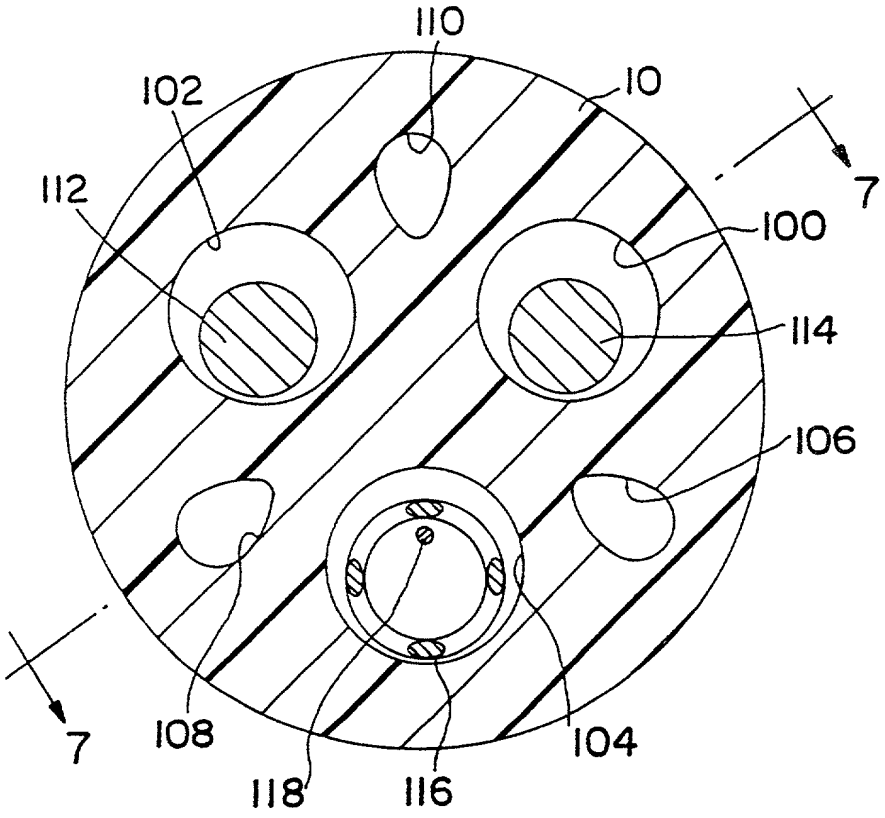
110
102
10
112
7
100
114
106
7
108
104
118
116

FIG. 3

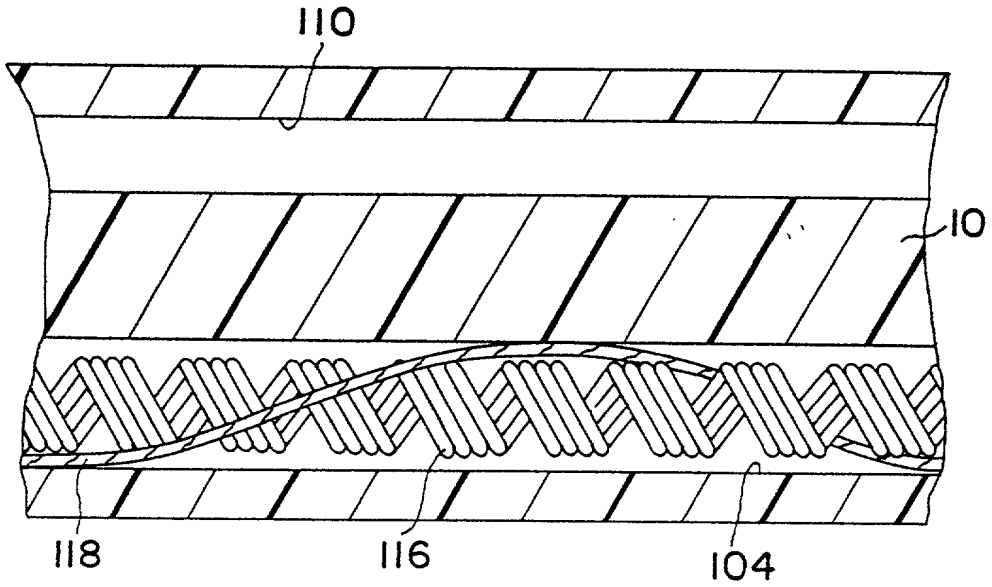

FIG. 4

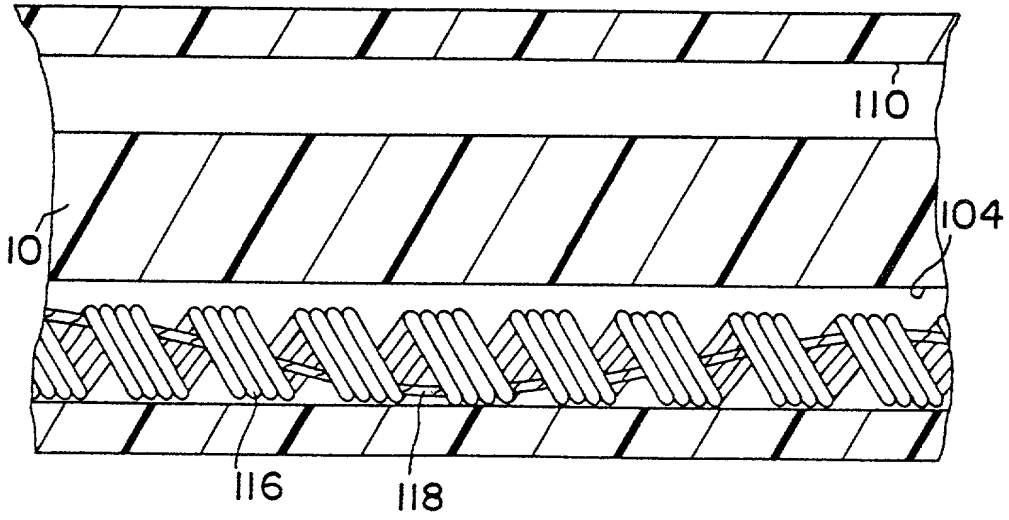

FIG. 5

D95-16592-07-400

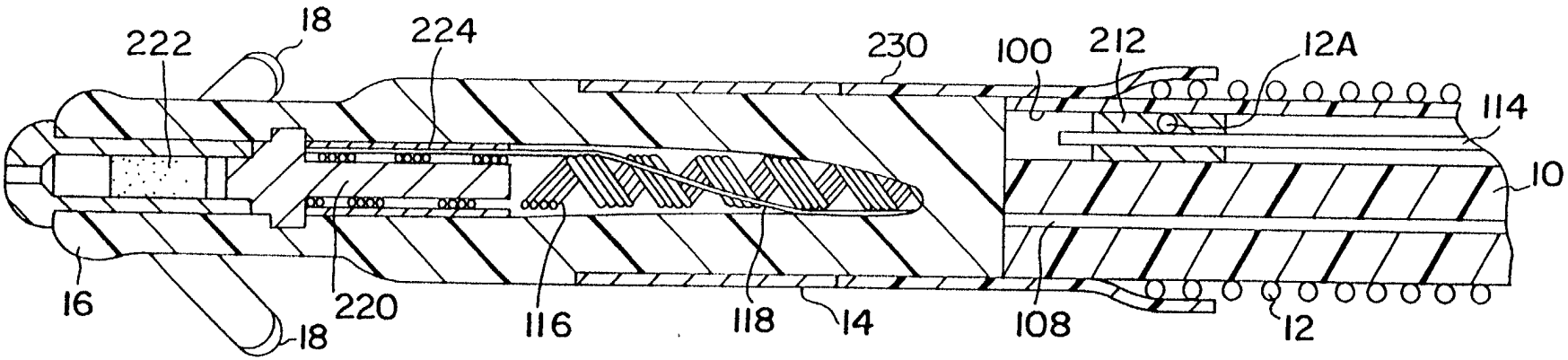
18
222
224
230
100
212
12A
114
10
16
220
116
118
14
108
12
18

FIG. 6

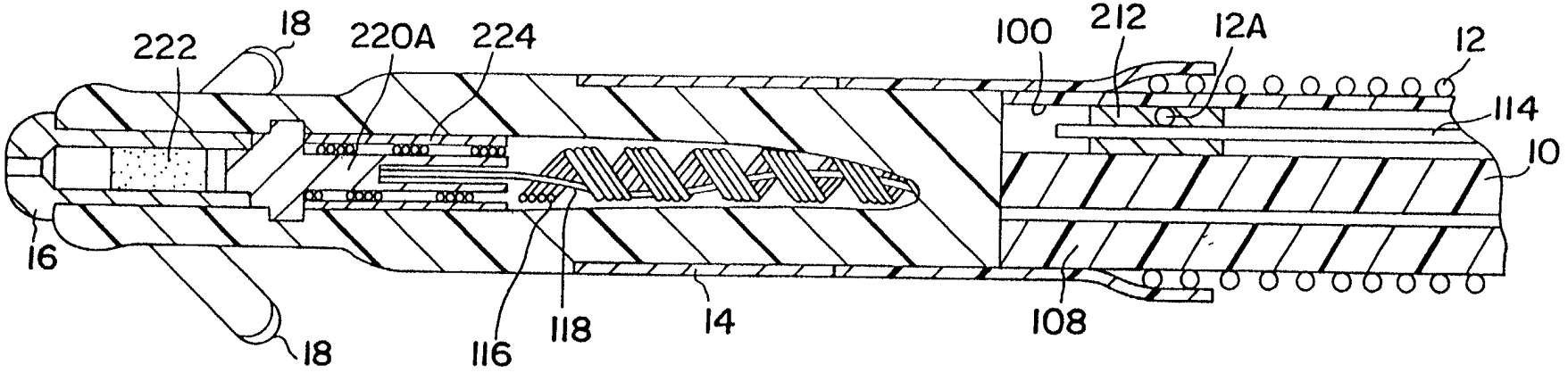
18
222
220A
224
100
212
12A
12
114
10
16
116
118
14
108
18

FIG. 7

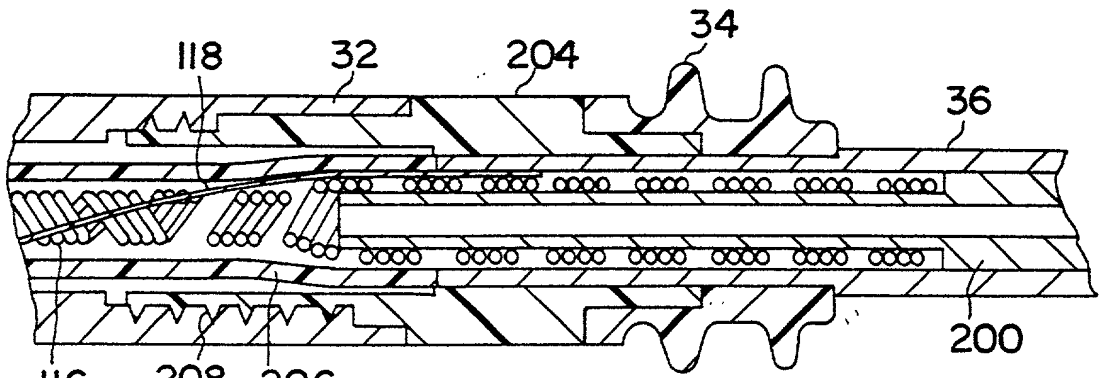

FIG. 8

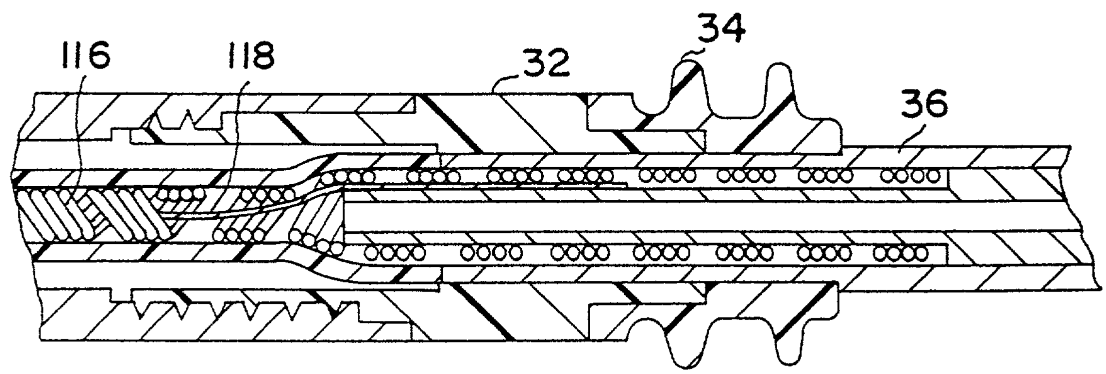

FIG. 9

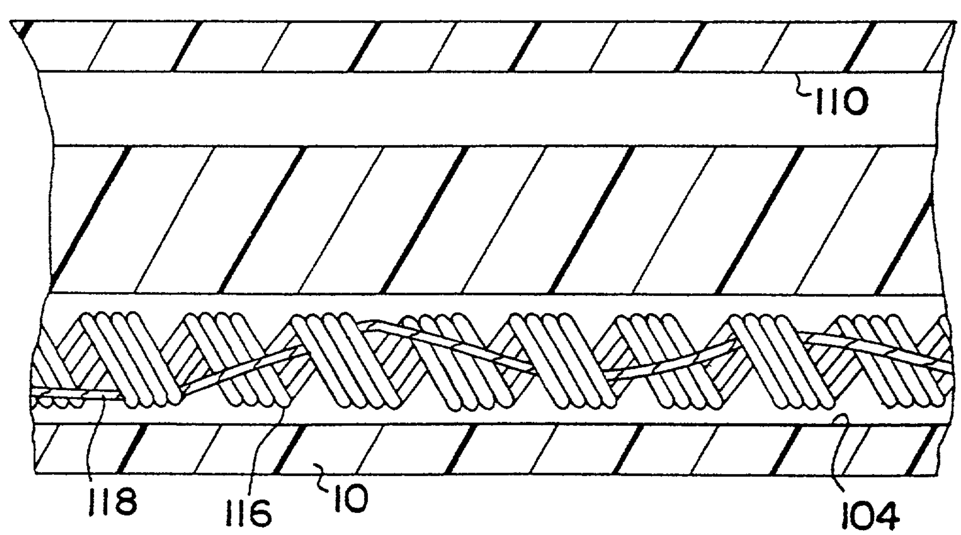

FIG. 10

0654592 • 07/1400

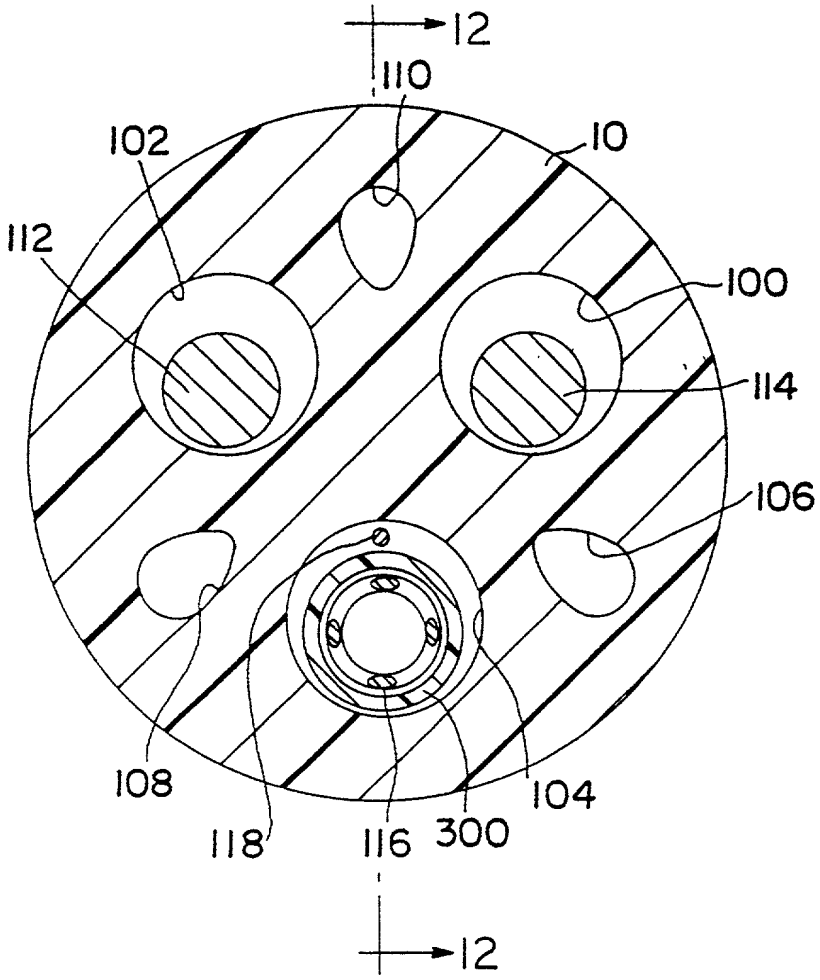

FIG. 11

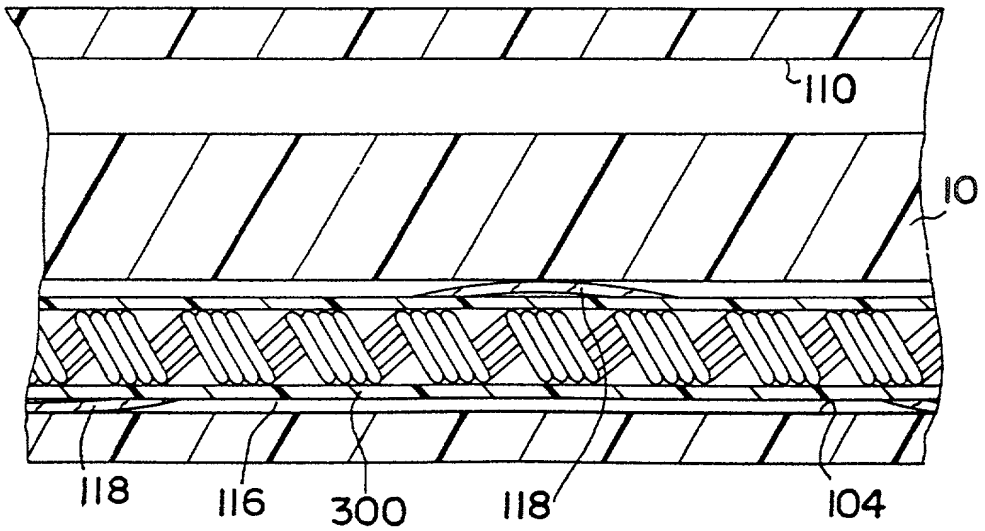

FIG. 12

0047400 09/17/92

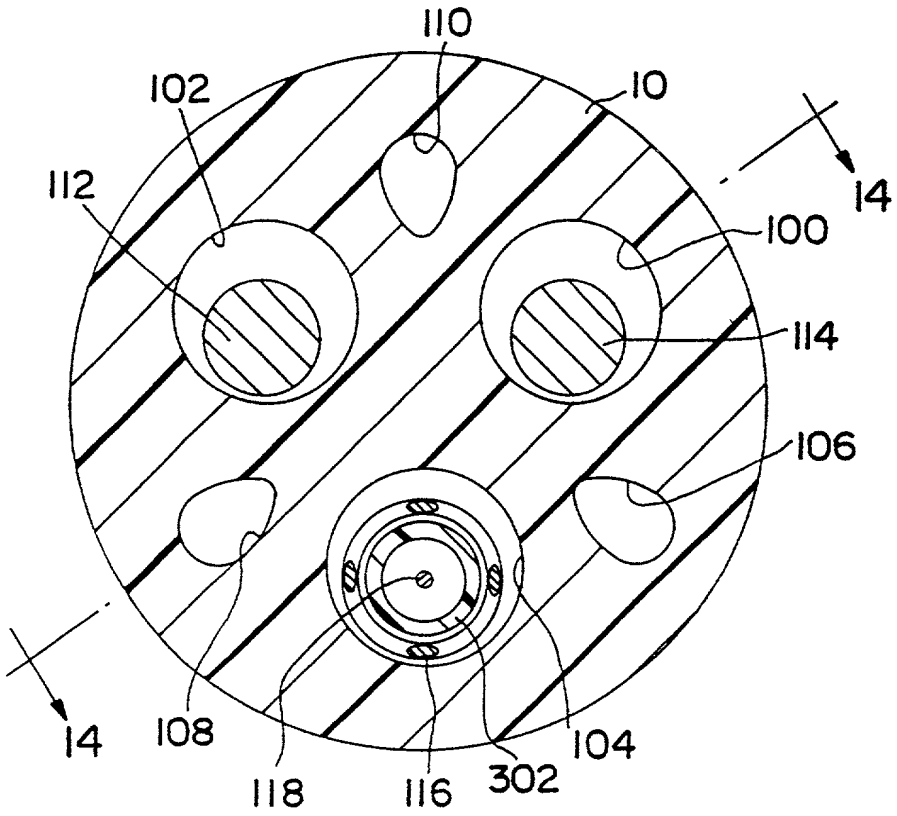

FIG. 13

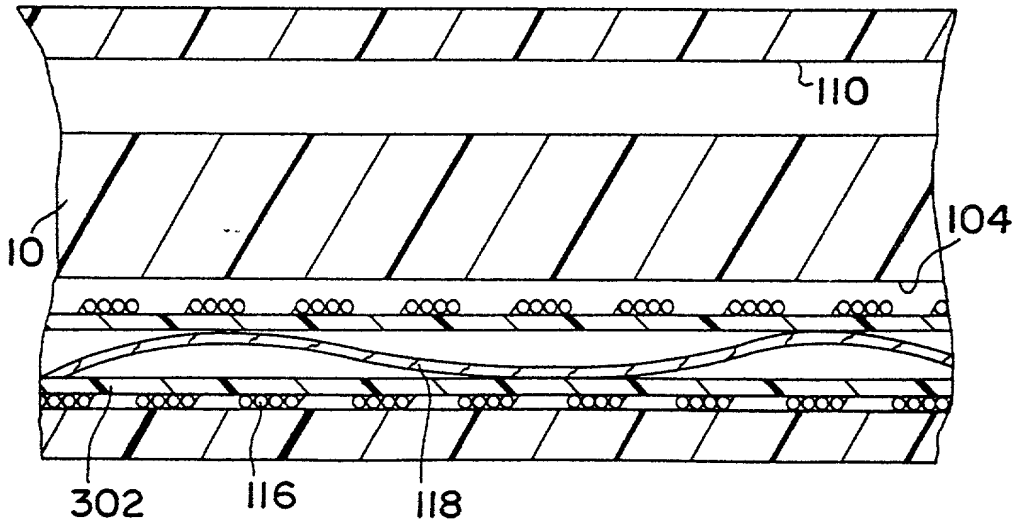

FIG. 14

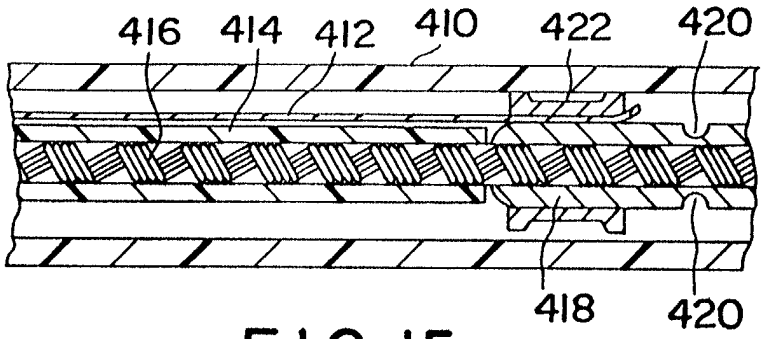
416
414
412
410
422
420
418
420

FIG. 15

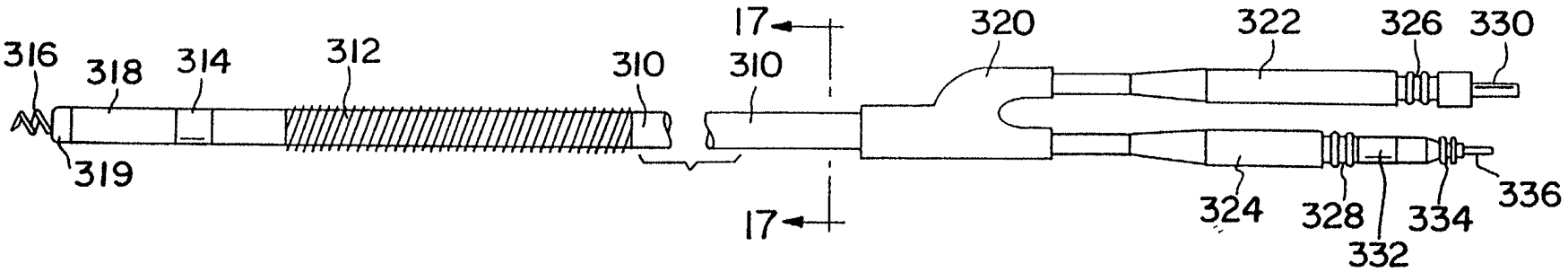
316
318
314
312
310
17
310
320
322
326
330
319
17
324
328
332
334
336

FIG. 16

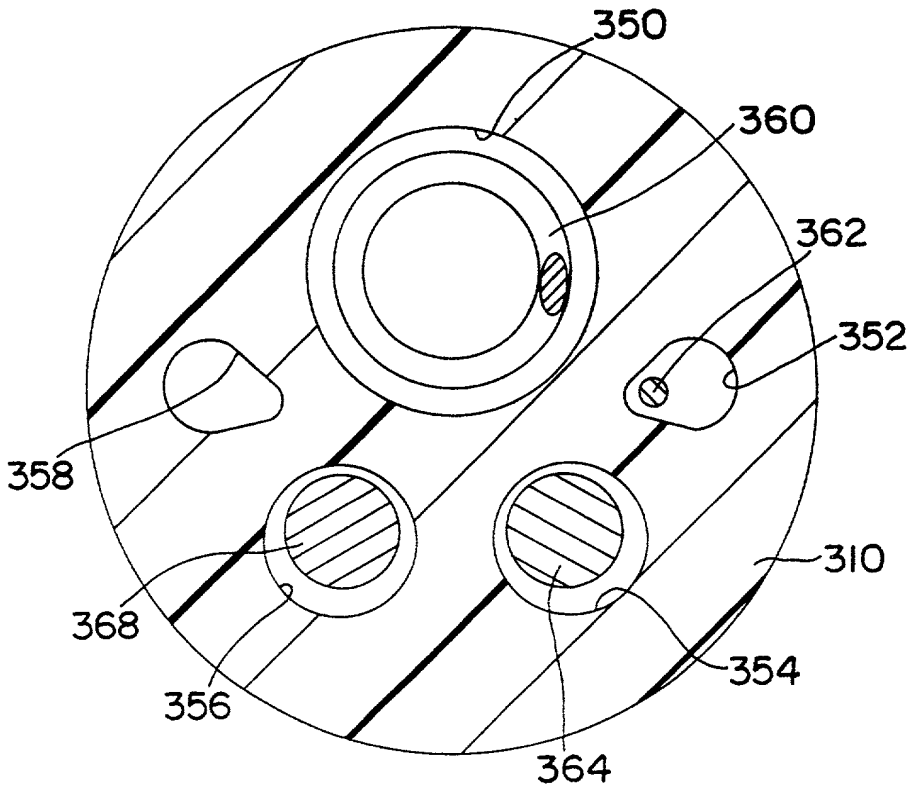
350
360
362
352
358
310
368
354
356
364

FIG. 17

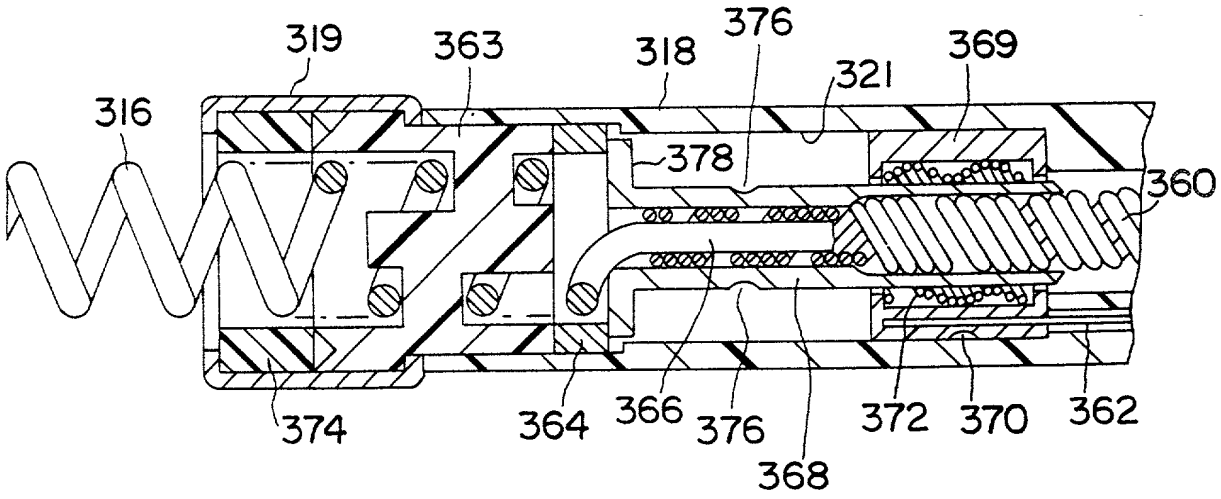
319
363
318
376
369
321
316
378
360
374
364
366
376
372
370
362
368

FIG. 18

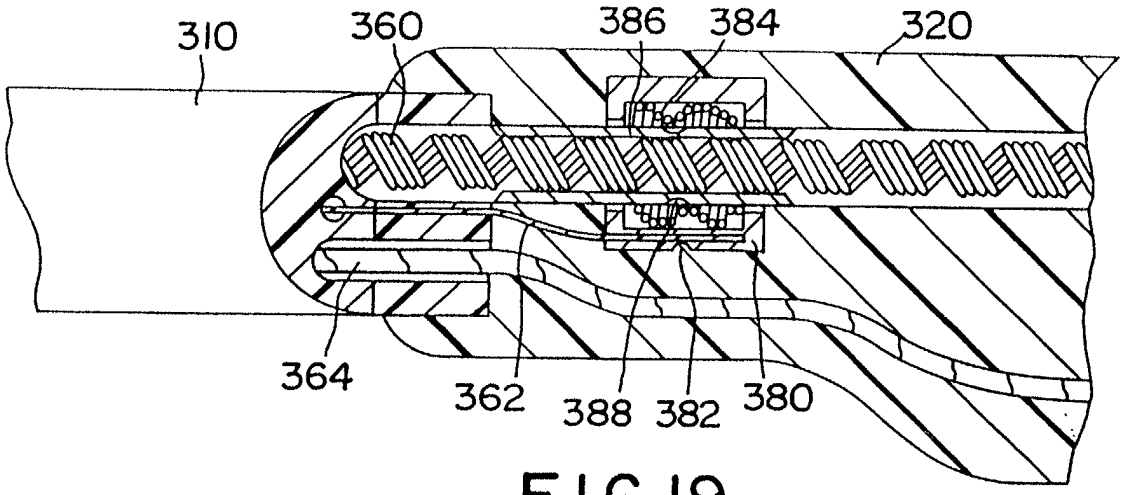

FIG. 19

TTORNEY DOCKET: P-4824

PATENT

United States Patent Application

COMBINED DECLARATION AND POWER OF ATTORNEY

As a below named inventor I hereby declare that: my residence, post office address and citizenship are as stated below ext to my name; that

I verily believe I am the original, first and sole inventor (if only one name is listed below) or a joint inventor (if ural inventors are named below) of the subject matter which is claimed and for which a patent is sought on the invention ntitled: MEDICAL ELECTRICAL LEAD.

e specification of which
X is attached hereto
_____ was filed on _____ application serial no. _____ was amended on ___ (if applicable) (in the case of a PCT-iled application) described and claimed in international no. ___ filed ___ and as amended on __ (if any), which I have reviewed nd for which I solicit a United States patent.

hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as ended by any amendment referred to above.

acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title , Code of Federal Regulations, §1.56(a).

hereby claim foreign priority benefits under Title 35, United States Code, §119/365 of any foreign application(s) for patent f inventor's certificate listed below and have also identified below any foreign application for patent or inventor's ertificate having a filing date before that of the application on the basis of which priority is claimed:

X no such applications have been filed.
___ such applications have been filed as follows:

FOREIGN APPLICATION(S), IF ANY, CLAIMING PRIORITY UNDER 35 USC §119

COUNTRY	APPLICATION NUMBER	DATE OF FILING	DATE OF ISSUE

ALL FOREIGN APPLICATIONS, IF ANY, FILED BEFORE THE PRIORITY APPLICATION(S)

COUNTRY	APPLICATION NUMBER	DATE OF FILING	DATE OF ISSUE

hereby claim the benefit under Title 35, United States Code, §120/365 of any United States and PCT international pplication(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the rior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, I cknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, §156(a) which ccurred between the filing date of the prior application and the national or PCT international filing date of this application.

---

1 § 1.56 Duty of disclosure: fraud, striking or rejection of applications.

(a) A duty of candor and good faith toward the Patent and Trademark Office rests on the inventor, on each attorney or agent who prepares or prosecutes the application and on every other individual who is substantively involved in the preparation or prosecution of the application and who is associated with the inventor, with the assignee or with anyone to whom there is an obligation to assign the application. All such individuals have a duty to disclose to the Office information they are aware of which is material to the examination of the application. Such information is material where there is substantial likelihood that a reasonable examiner would consider it important in deciding whether to allow the application to issue as a patent. The duty is commensurate with the degree of involvement in the preparation or prosecution of the application.

U.S. APPLICATION NUMBER	DATE OF FILING	STATUS (patented, pending, abandoned)
60/033,611	19 December 1996	pending

I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected herewith:

Harold R. Patton	Reg. No. 22,157	Dianne M.F. Plunkett	Reg. No. 35,649
Reed A. Duthler	Reg. No. 30,625	Michael J. Jaro	Reg. No. 34,472
Daniel W. Latham	Reg. No. 30,401	Curtis D. Kinghorn	Reg. No. 33,926
Michael B. Atlas	Reg. No. 30,606	Thomas F. Woods	Reg. No. 36,725
		Peter Forrest	Reg. No. 33,235

Please direct all correspondence in this case to: Medtronic, Inc.
7000 Central Avenue N.E, Minneapolis, Minnesota 55432
Telephone No. (612) 574-3156

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

	Full Name of Inventor	FIRST NAME David	MIDDLE INITIAL D.	LAST NAME Vesness
201	Residence & Citizenship	CITY Forest Lake	STATE OR FOREIGN COUNTRY Minnesota	COUNTRY of CITIZENSHIP US
	Post Office Address	POST OFFICE ADDRESS 909 12th Avenue Southeast	CITY Forest Lake	STATE/ZIP/COUNTRY MN/55025/US
SIGNATURE OF INVENTOR 201: David D Vesness				DATE April 16, 1997
	Full Name of Inventor	FIRST NAME George	MIDDLE INITIAL M.	LAST NAME Huepenbecker
202	Residence & Citizenship	CITY Vadnais Heights	STATE OR FOREIGN COUNTRY Minnesota	COUNTRY OF CITIZENSHIP US
	Post Office Address	POST OFFICE ADDRESS 3450 Montmorency	CITY Vadnais Heights	STATE/ZIP/COUNTRY MN/55110/US
SIGNATURE OF INVENTOR 202: George M. Huepenbecker				DATE April 16, 1997
	Full Name of Inventor	FIRST NAME Dale	MIDDLE NAME A.	LAST NAME Wahlstrom
203	Residence & Citizenship	CITY Plymouth	STATE OR FOREIGN COUNTRY Minnesota	COUNTRY OF CITIZENSHIP US
	Post Office Address	POST OFFICE ADDRESS 4585 Goldenrod Lane	CITY Plymouth	STATE/ZIP/COUNTRY MN/55442/US
SIGNATURE OF INVENTOR 203: Dale A Wahlstrom				DATE 4/16/97

X THIS IS THE FINAL PAGE OF THIS DECLARATION.

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: VERNESS et al

Examiner: ---.

Serial # : ---

Group Art Unit: ---

Filed : HEREWITH

Docket: P-4824.06 CON

Title: MEDICAL ELECTRICAL LEAD

ASSOCIATE POWER OF ATTORNEY

Commissioner of Patents
and Trademarks
Washington, D.C. 20231

Sir:

Please recognize Girma Wolde-Michael, Reg. No. 36,724, Eric R. Waldkoetter, Reg. No. 36,713, Beth L. McMahon, Reg. No. 41,987, and Kenneth J. Collier as associate attorneys for the above-identified patent application and having full authority to prosecute all matters in the Patent and Trademark Office. Please direct all further correspondence to: GIRMA WOLDE-MICHAEL, Medtronic, Inc., 7000 Central Avenue N.E., Minneapolis, Minnesota 55432.

Respectfully submitted,

7/14/00
Date

Daniel W. Latham
Reg. No. 30,401

DOCKET NO. P-4824.06 CON